

Document made available under the Patent Cooperation Treaty (PCT)

International application number: PCT/GB04/005392

International filing date: 20 December 2004 (20.12.2004)

Document type: Certified copy of priority document

Document details: Country/Office: GB
Number: 0329364.4
Filing date: 19 December 2003 (19.12.2003)

Date of receipt at the International Bureau: 24 January 2005 (24.01.2005)

Remark: Priority document submitted or transmitted to the International Bureau in compliance with Rule 17.1(a) or (b)



World Intellectual Property Organization (WIPO) - Geneva, Switzerland
Organisation Mondiale de la Propriété Intellectuelle (OMPI) - Genève, Suisse



INVESTOR IN PEOPLE

The Patent Office
Concept House
Cardiff Road
Newport
South Wales
NP10 8QQ

I, the undersigned, being an officer duly authorised in accordance with Section 74(1) and (4) of the Deregulation & Contracting Out Act 1994, to sign and issue certificates on behalf of the Comptroller-General, hereby certify that annexed hereto is a true copy of the documents as originally filed in connection with the patent application identified therein.

In accordance with the Patents (Companies Re-registration) Rules 1982, if a company named in this certificate and any accompanying documents has re-registered under the Companies Act 1980 with the same name as that with which it was registered immediately before re-registration save for the substitution as, or inclusion as, the last part of the name of the words "public limited company" or their equivalents in Welsh, references to the name of the company in this certificate and any accompanying documents shall be treated as references to the name with which it is so re-registered.

In accordance with the rules, the words "public limited company" may be replaced by p.l.c., plc, P.L.C. or PLC.

Re-registration under the Companies Act does not constitute a new legal entity but merely subjects the company to certain additional company law rules.

Signed

Dated 13 January 2005

Patents Form 1/77

Patents Act 1977
(Rule 16)

THE PATENT OFFICE
PE
19 DEC 2003
RECEIVED BY FAX



1/77

Request for grant of a patent

(See the notes on the back of this form. You can also get an explanatory leaflet from the Patent Office to help you fill in this form)

19 DEC 2003

The Patent Office

Cardiff Road
Newport
South Wales
NP10 8QQ

1. Your reference

CDT499 GB

19DEC03 EB60591-1 D10063

P01/7700 0.00-0329364.4 ACCOUNT CHA

2. Patent application number

(The Patent Office will fill in this part)

0329364.4

3. Full name, address and postcode of the or of each applicant (*underline all surnames*)

CAMBRIDGE DISPLAY TECHNOLOGY LIMITED

Greenwich House
Madingley Rise
Madingley Road
Cambridge CB3 0TX

Patents ADP number (*if you know it*)

If the applicant is a corporate body, give the country/state of its incorporation

United Kingdom

06166441004

4. Title of the invention

OPTICAL DEVICE

5. Name of your agent (*if you have one*)

"Address for service" in the United Kingdom to which all correspondence should be sent (*including the postcode*)

Cambridge Display Technology Limited
c/o IP Department
Greenwich House
Madingley Rise
Madingley Road
Cambridge
CB3 0TX

Patents ADP number (*if you know it*)

06166441004

6. If you are declaring priority from one or more earlier patent applications, give the country and the date of filing of the or of each of these earlier applications and (*if you know it*) the or each application number

Country

Priority application number
(*if you know it*)Date of filing
(*day / month / year*)

7. If this application is divided or otherwise derived from an earlier UK application, give the number and the filing date of the earlier application

Number of earlier application

Date of filing
(*day / month / year*)

8. Is a statement of inventorship and of right to grant of a patent required in support of this request? (*Answer 'Yes' if:*

YES

a) any applicant named in part 3 is not an inventor, or
b) there is an inventor who is not named as an applicant, or

c) any named applicant is a corporate body.
See note (d))

Patents Form 1/77

0089411 19 Dec 03 10:38

Patents Form 1/77


9. Enter the number of sheets for any of the following items you are filing with this form. Do not count copies of the same document

Continuation sheets of this form

Description 19

Claim(s) 3

Abstract -

Drawing(s) 1 

10. If you are also filing any of the following, state how many against each item.

Priority documents -

Translations of priority documents -

Statement of inventorship and right to grant of a patent (Patents Form 7/77) -

Request for preliminary examination and search (Patents Form 9/77) 1

Request for substantive examination (Patents Form 10/77) 1

Any other documents (please specify)

11. I/We request the grant of a patent on the basis of this application.

Signature

Date

18 December 2003

12. Name and daytime telephone number of person to contact in the United Kingdom

Anwar Gillani
Tel: 01223 723519**Warning**

After an application for a patent has been filed, the Comptroller of the Patent Office will consider whether publication or communication of the invention should be prohibited or restricted under Section 22 of the Patents Act 1977. You will be informed if it is necessary to prohibit or restrict your invention in this way. Furthermore, if you live in the United Kingdom, Section 23 of the Patents Act 1977 stops you from applying for a patent abroad without first getting written permission from the Patent Office unless an application has been filed at least 6 weeks beforehand in the United Kingdom for a patent for the same invention and either no direction prohibiting publication or communication has been given, or any such direction has been revoked.

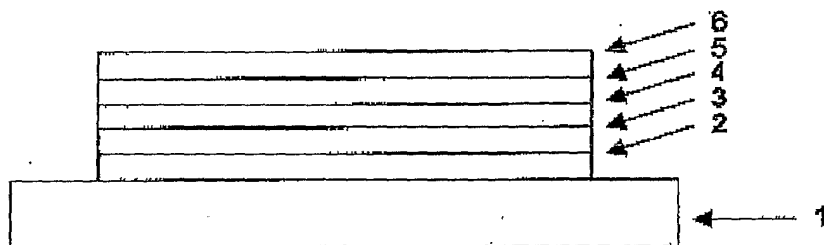
Notes

- If you need help to fill in this form or you have any questions, please contact the Patent Office on 08459 500505.
- Write your answers in capital letters using black ink or you may type them.
- If there is not enough space for all the relevant details on any part of this form, please continue on a separate sheet of paper and write "see continuation sheet" in the relevant part(s). Any continuation sheet should be attached to this form.
- If you have answered 'Yes' Patents Form 7/77 will need to be filed.
- Once you have filled in the form you must remember to sign and date it.
- For details of the fee and ways to pay please contact the Patent Office.

Patents Form 1/77

0089411 19-Dec-03 10:38

Figure 1



DUPLICATE

Optical device

Field of the Invention

This invention relates to organic optical devices comprising a layer of insoluble organic material and methods for the production thereof.

Background of the Invention

Organic semiconductors are now frequently used in a number of optical devices such as in organic light emitting diodes ("OLEDs") as disclosed in WO 90/13148, photovoltaic devices as disclosed in WO 96/16449 and photodetectors as disclosed in US 5523555.

A typical OLED comprises a substrate, on which is supported an anode (commonly indium tin oxide or "ITO"), a cathode and an organic electroluminescent layer between the anode and cathode. In operation, holes are injected into the device through the anode and electrons are injected into the device through the cathode. The holes and electrons combine in the organic electroluminescent layer to form an exciton which then undergoes radiative decay to give light. Other layers may be present in the OLED, for example a layer of organic hole injection material such as poly(ethylene dioxy thiophene) / polystyrene sulfonate (PEDT / PSS) may be provided between the anode and the organic electroluminescent layer to assist injection of holes from the anode to the organic electroluminescent layer.

Various classes of organic light emitting materials are known, in particular: polymers such as poly(p-phenylenevinylene) (as disclosed in WO 90/13148), polyfluorenes and polyphenylenes; the class of materials known as small molecule materials such as tris-(8-hydroxyquinoline)aluminium ("Alq₃") as disclosed in US 4,539,507; and the class of materials known as dendrimers as disclosed in WO 99/21935. These materials electroluminesce by radiative decay of singlet excitons (i.e. fluorescence) however spin statistics dictate that up to 75% of excitons are triplet excitons which undergo non-radiative decay, i.e. quantum efficiency may be as low as 25% for fluorescent OLEDs – see, for example, *Chem. Phys. Lett.*, 1993, 210, 61, *Nature (London)*, 2001, 409, 494, *Synth. Met.*, 2002, 125, 55 and references therein.

Accordingly, considerable effort has been directed towards producing luminescence from triplet excitons (phosphorescence) by utilising spin-orbit coupling effects in metal complexes that enable triplet excitons to undergo radiative decay. The metal complex is doped into a host material from which it receives charge and / or triplet excitons. Examples of complexes investigated for this purpose include lanthanide metal chelates [*Adv. Mater.*, 1999, 11, 1349], a

platinum (II) porphyrin [*Nature (London)*, 1998, 395, 151] and *tris*(phenylpyridine) iridium (III) (hereinafter "Ir(ppy)₃") [*Appl. Phys. Lett.*, 1999, 75, 4; *Appl. Phys. Lett.*, 2000, 77, 904]. Fuller reviews of such complexes may be found in *Pure Appl. Chem.*, 1999, 71, 2095, *Materials Science & Engineering, R: Reports* (2002), R39(5-6), 143-222 and *Polymeric Materials Science and Engineering* (2000), 83, 202-203.

Prior art phosphorescent OLEDs often comprise charge transporting and / or charge blocking layers used in conjunction with the electroluminescent layer in order to maximise device efficiency. The charge transporting / blocking layers and the electroluminescent layers are typically formed by vacuum evaporation of the appropriate materials in sequence.

Deposition of materials from solution, for example by spin-coating or inkjet printing, offers advantages over vacuum deposition such as simplified processing. However, solution deposition of multiple layers is complicated by the tendency of initially cast or deposited layers to dissolve in the solvents used for succeeding layers. Thus, solution deposition will commonly be employed for one layer only. For example, US 2002/096995 discloses spin-coating of an electroluminescent layer of polyvinylcarbazole (hereinafter "PVK") host doped with Ir(ppy)₃ emitter onto a substrate of ITO and PEDT / PSS. A layer of electron transporting material is then deposited by vacuum evaporation.

One solution to this problem is disclosed in JP 2003-077673 wherein a hole transporting layer of PVK is formed by spin-coating from 1,2-dichloroethane solution followed by formation of an electroluminescent layer of 9,9-dioctylfluorene host / Ir(ppy)₃ emitter by spin-coating from xylene solution. The formation of this bilayer by solution processing is possible due to the low solubility of PVK in the xylene solvent used for the electroluminescent layer. This approach is limited in that the material used for the first layer can only be selected from those that do not dissolve in the solvent used for the second layer.

Another solution to this problem is disclosed in JP 2002-050482 wherein a first layer of insoluble poly(phenylvinylene) (hereinafter "PPV") is formed by deposition by spin-coating of a soluble precursor compound, followed by thermal conversion of the precursor to insoluble PPV. A layer of PVK host / Ir(ppy)₃ guest is then deposited onto the insoluble PPV layer. Again, this approach is limited in that it is only applicable to insoluble compounds that have a soluble precursor form. Furthermore, the thermal conversion required by these precursors requires forcing conditions and generates corrosive by-products that may harm the performance of the finished device.

It is therefore an object of the invention to provide a method of forming a phosphorescent OLED by solution deposition of a plurality of layers that is not constrained by the nature of the material being used. It is a further object of the invention to provide a phosphorescent OLED having improved device performance.

Summary of the invention

The present inventors have found that a phosphorescent OLED comprising a plurality of solution processable layers may be formed by treatment of a solution processable layer to render it insoluble.

Accordingly, in a first aspect the invention provides a method of forming an organic light emitting diode comprising the steps of:

- providing a substrate comprising a first electrode for injection of charge carriers of a first type
- forming a charge transporting layer by depositing onto the substrate a charge transporting material for transporting charge carriers of the first type, the charge transporting material being soluble in a solvent;
- treatment of the charge transporting layer to render it insoluble in the solvent;
- forming an electroluminescent layer by depositing onto the charge transporting layer a composition comprising the solvent, a phosphorescent material and a host material; and
- depositing onto the electroluminescent layer a second electrode for injection of charge carriers of a second type.

Preferably, the first electrode is an anode; the second electrode is a cathode; the charge carriers of the first type are holes; and the charge carriers of the second type are electrons.

Preferably, the charge transporting material is deposited from solution. The solvent used for such a solution may be the same as or different to the solvent used to deposit the electroluminescent layer.

In one embodiment, the charge transporting material comprises a cross-linkable material and the treatment comprises subjecting the charge transporting layer to heat or electromagnetic radiation, in particular UV radiation, in order to cross-link the charge transporting material.

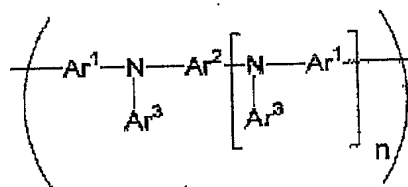
The present inventors have surprisingly found that the charge transporting layer may be rendered insoluble in the solvent even when cross-linking groups are absent. Accordingly, in another embodiment the charge transporting layer is substantially free of cross-linkable groups and the treatment comprises subjecting the charge transporting layer to heat.

Following the aforementioned treatment, it may be possible to solubilise the charge transporting layer under forcing conditions (e.g. by exposure to the solvent at elevated temperature and / or immersion in the solvent over a prolonged period). However, the present inventors have found that the charge transporting layer remains intact under conditions typically employed for deposition of the electroluminescent layer, and the term "insoluble" should be construed accordingly.

Preferably, the charge transporting material is a polymer.

Preferably, the polymer comprises an optionally substituted triarylamine repeat unit.

Preferably, the triarylamine repeat unit comprises an optionally substituted first repeat unit of formula (I):



(I)

wherein each Ar^1 , Ar^2 and Ar^3 is the same or different and independently represents optionally substituted aryl or heteroaryl; and n is 0 or 1.

Preferably, Ar^1 is phenylene.

Preferably, Ar^2 is 1,4-phenylene or 4,4'-biphenylene. When n is 0, Ar^2 is preferably 1,4-phenylene. When n is 1, Ar^2 is preferably 1,4-phenylene or 4,4'-biphenylene.

Preferably, Ar^3 is phenyl. Ar^3 may be substituted or unsubstituted; preferably Ar^3 is substituted with a solubilising group, more preferably optionally substituted C_{1-20} alkyl or C_{1-20} alkoxy.

Preferably, the polymer is a copolymer comprising two or more repeat units. Preferably, one of the repeat units of such a copolymer is the first repeat unit of formula (I). More preferably, the

copolymer comprises a second repeat unit selected from optionally substituted fluorene, indenofluorene, spirofluorene and phenylene.

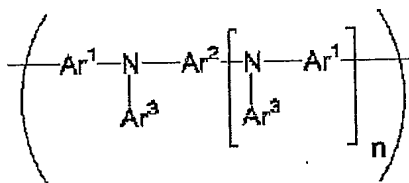
Preferably, the phosphorescent material is a metal complex.

Preferably, the host material is a host polymer.

Preferably, the host polymer is a polymer comprising a triarylamine repeat unit as defined above, preferably a first repeat unit.

In a second aspect, the invention provides an organic light emitting diode obtainable by the method according to the first aspect of the invention.

In a second aspect, the invention provides an organic light emitting diode comprising, in sequence, an anode; a hole transporting layer; an electroluminescent layer comprising a phosphorescent material and a host material; and a cathode, wherein the hole transporting layer is a polymer comprising an optionally substituted repeat unit of formula (I):



(I)

wherein each Ar^1 , Ar^2 and Ar^3 is the same or different and independently represents optionally substituted aryl; and n is 0 or 1.

Preferably, the polymer comprises a repeat unit selected from optionally substituted fluorene, indenofluorene, spirofluorene and phenylene.

Preferably, a hole injecting layer comprising a conductive organic material is located between the anode and the hole transporting layer.

Preferably, the phosphorescent material is a metal complex.

Brief Description of the Drawings

The present invention will now be described in further detail, by way of example only, with reference to the accompanying drawings in which:

FIGURE 1 shows an OLED prepared according to the method of the invention.

Detailed Description of the Invention

With reference to Figure 1, an organic light emitting diode prepared according to the method of the invention comprises a substrate 1, an anode 2 of indium tin oxide, a layer 3 of organic hole injection material, an insoluble hole transporting layer 4, an electroluminescent layer 5 and a cathode 6.

Optical devices tend to be sensitive to moisture and oxygen. Accordingly, the substrate 1 preferably has good barrier properties for prevention of ingress of moisture and oxygen into the device. The substrate is commonly glass, however alternative substrates may be used, in particular where flexibility of the device is desirable. For example, the substrate may comprise a plastic as in US 6268695 which discloses a substrate of alternating plastic and barrier layers or a laminate of thin glass and plastic as disclosed in EP 0949850.

Although not essential, the presence of layer 3 of organic hole injection material is desirable as it assists hole injection from the anode into the layer or layers of semiconducting polymer. Examples of organic hole injection materials include PEDT / PSS as disclosed in EP 0901176 and EP 0947123, or polyaniline as disclosed in US 5723873 and US 5798170.

Cathode 6 is selected in order that electrons are efficiently injected into the device and as such may comprise a single conductive material such as a layer of aluminium. Alternatively, it may comprise a plurality of metals, for example a bilayer of calcium and aluminium as disclosed in WO 98/10621. A thin layer of dielectric material such as lithium fluoride may be provided between the electroluminescent layer 5 and the cathode 6 to assist electron injection as disclosed in, for example, WO 00/48258.

The device is preferably encapsulated with an encapsulant (not shown) to prevent ingress of moisture and oxygen. Suitable encapsulants include a sheet of glass, films having suitable barrier properties such as alternating stacks of polymer and dielectric as disclosed in, for example, WO 01/81649 or an airtight container, optionally with a desiccant, as disclosed in, for example, WO 01/19142.

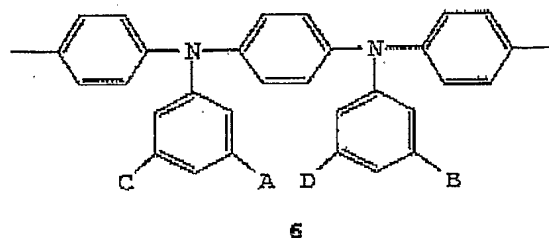
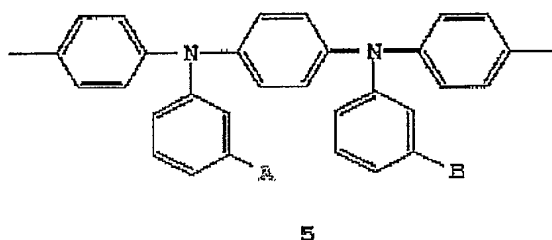
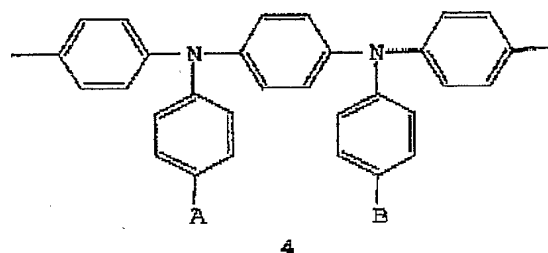
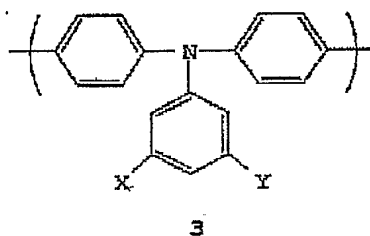
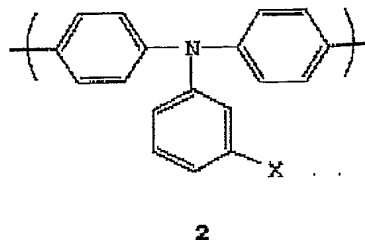
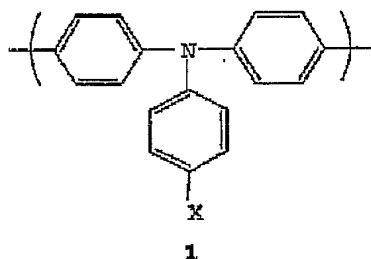
In a practical device, at least one of the electrodes is semi-transparent in order that light may be emitted. Where the anode is transparent, it typically comprises indium tin oxide. Examples of transparent cathodes are disclosed in, for example, GB 2348316.

A typical electroluminescent device comprises an anode having a workfunction of 4.8 eV. Accordingly, the HOMO level of the hole transporting material for hole transporting layer 4 is preferably around 4.8-5.5 eV. Suitable hole transporting materials are small molecules, polymers and dendrimers comprising optionally substituted triarylamine units, in particular triphenylamine units, and materials comprising carbazole units.

The hole transporting layer 4 may also serve as an electron blocking layer. Electron blocking functionality is provided if the hole transporting layer 4 possesses a shallower (i.e. less positive) LUMO level(s) than the host material and the phosphorescent dopant. For a typical device architecture, suitable LUMO level for the hole transporting layer 4 is about 1.6-2.3eV.

The hole transporting layer 4 may also serve as an exciton blocking layer. Exciton blocking functionality is provided if the hole transporting layer 4 has (a) a wider HOMO-LUMO bandgap and (b) a wider T_1 - T_0 energy gap than the phosphorescent dopant.

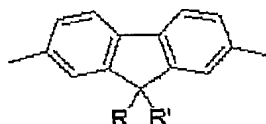
Preferred hole transporting materials for hole transporting layer 4 are polymers comprising triarylamine repeat units such as polymers having repeat units 1-6:



wherein X, Y, A, B, C and D are independently selected from H or a substituent group. More preferably, one or more of X, Y, A, B, C and D is independently selected from the group consisting of optionally substituted, branched or linear alkyl, aryl, perfluoroalkyl, thioalkyl, cyano, alkoxy, heteroaryl, alkylaryl and arylalkyl groups. Most preferably, X, Y, A and B are C₁₋₁₀ alkyl. The repeat unit of formula 1 is most preferred.

These polymers may be homopolymers or copolymers. Where they are copolymers, suitable co-repeat units are optionally substituted arylenes such as fluorenes as disclosed in Adv. Mater. 2000 12(23) 1737-1750, particularly 2,7-linked 9,9-dialkyl fluorene, 9,9-diaryl fluorene or 9-alkyl-9-aryl fluorene; a spirofluorene such as 2,7-linked spirobifluorene as disclosed in EP 0707020; an indenofluorene such as a 2,7-linked indenofluorene; or a phenylene such as alkyl or alkoxy substituted 1,4-phenylene. Each of these repeat units may be substituted. These copolymers are particularly advantageous because they have good film forming properties and may be readily formed by Suzuki or Yamamoto polymerisation which enables a high degree of control over the regioregularity of the resultant polymer.

Preferred polyfluorenes comprise repeat units of formula (II):



(II)

wherein R and R' are independently selected from hydrogen or optionally substituted alkyl, alkoxy, aryl, arylalkyl, heteroaryl and heteroarylalkyl, and at least one of R and R' is not hydrogen. More preferably, at least one of R and R' comprises an optionally substituted C₁-C₂₀ alkyl group, more preferably C₄-C₁₀ alkyl group.

Preferably, the first semiconducting polymer comprises a 1:1 regular, alternating copolymer of a fluorene repeat unit and a triarylamine repeat unit.

Another suitable hole transporting polymer is poly(vinyl carbazole).

The hole transporting material is soluble in at least some organic solvents, in particular to allow its deposition from solution. Following deposition, the hole transporting material is treated so as to render it insoluble.

One suitable treatment entails cross-linking of cross-linkable groups that are either bound to or blended with the hole transporting material. Suitable cross-linkable groups include oxetanes as disclosed in Nature 421, 829-833, 2003; azides; acrylates as disclosed in WO 96/20253; vinyl groups as disclosed in US 6107452; and ethynyl groups as disclosed in Kim *et al*, Synthetic Metals 122 (2001), 363-368. Cross-linking may be achieved through thermal treatment or exposure of the hole transporting layer to radiation, in particular UV radiation.

The hole transporting layer may be rendered insoluble even if no cross-linkable groups are present within it by an appropriate insolubilisation treatment. The hole transporting layer is then stable to solution deposition of the electroluminescent layer. The present inventors have found that this insoluble layer forms (a) regardless of whether a PEDT/PSS layer is used or not (although presence of PEDT/PSS is preferred) and (b) in air or in a nitrogen only environment. However, the present inventors have found that it is necessary to subject the hole transporting layer to an insolubilising treatment in order to maximise device performance, and in particular to minimise mixing of the second layer with the first layer by increasing the insolubility of the first layer. Without wishing to be bound by any theory, possible mechanisms for loss of solubility of the hole transporting layer upon insolubilisation treatment when no cross-linkable groups are present include formation of a relatively solvent impermeable surface or an adhesion to the

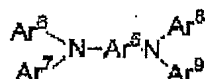
surface that the hole transporting layer following elimination of solvent from a solution deposited hole transporting layer.

Means for eliminating solvent include vacuum treatment and / or heat treatment. Heat treatment, either alone or in combination with vacuum treatment, is preferred. For heat treatment, a suitable treatment period is in the range 5 minutes - 2 hours. The heat treatment temperature is preferably above the glass transition temperature of the hole transporting material.

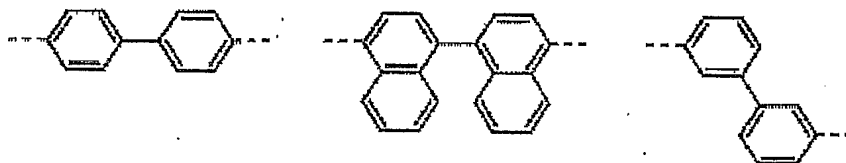
Preferably, the hole transporting layer has a thickness in the range 10-20 nm. If a hole transporting polymer is used to form the hole transporting layer 4, its thickness may be modified by appropriate selection of molecular weight (Mw) of the polymer. Thus, Mw of 50,000 may give a thickness as low as about 2 nm, but up to about 15-20 nm for Mw of around 250,000-300,000.

The electroluminescent layer 5 is deposited directly onto the hole transporting layer 4 and comprises a host material and a phosphorescent dopant. It is necessary for the host to possess a higher T_1 energy level than the dopant. Examples of suitable host materials are small molecules comprising triarylamine units (for examples see Shirota, J. Mater. Chem., 2000, 10, 1-26); polymers comprising triarylamine repeat units as described above; or carbazole units, in particular poly(vinylcarbazole).

The host material may also have charge transporting properties. Hole transporting host materials are particularly preferred such as the optionally substituted hole-transporting arylamine having the following formula:

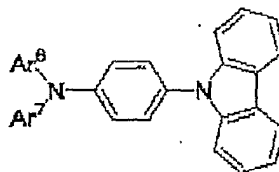


wherein Ar^5 is an optionally substituted aromatic group, such as phenyl, or



and Ar^6 , Ar^7 , Ar^8 and Ar^9 are optionally substituted aromatic or heteroaromatic groups (Shi et al (Kodak) US 5,554,450. Van Slyke et al, US 5,061,569. So et al (Motorola) US 5,853,905 (1997)). Ar is preferably biphenyl. At least two of Ar^6 , Ar^7 , Ar^8 and Ar^9 may be bonded to either a thiol group, or a group containing a reactive unsaturated carbon-carbon bond. Ar^6 and Ar^7

and/or Ar⁶ and Ar⁹ are optionally linked to form a N containing ring, for example so that the N forms part of a carbazole unit e.g.



Host materials may alternatively possess electron transporting properties. Examples of electron transporting host materials are azoles, diazoles, triazoles, oxadiazoles, benzoxazoles, benzazoles and phenanthrolines, each of which may optionally be substituted. Particularly preferred substituents are aryl groups, in particular phenyl. oxadiazoles, in particular aryl-substituted oxadiazoles. These host materials may exist in small molecule form or may be provided as repeat units of a polymer, in particular as repeat units located in the backbone of a polymer or as substituents pendant from a polymer backbone. Specific examples of electron transporting host materials include 3-phenyl-4-(1-naphthyl)-5-phenyl-1,2,4-triazole and 2, 9-dimethyl-4,7-diphenyl-phenanthroline.

Host materials may be bipolar, i.e. capable of transporting holes and electrons. Suitable bipolar materials preferably contain at least two carbazole units (Shirota, J. Mater. Chem., 2000, 10, 1-25). In one preferred compound, both Ar⁶ and Ar⁷ and Ar⁸ and Ar⁹ as described above are linked to form carbazole rings and Ar⁵ is phenyl. Alternatively, a bipolar host material may be a material comprising a hole transporting segment and an electron transporting segment. An example of such a material is a polymer comprising a hole transporting segment and an electron transporting segment as disclosed in WO 00/55927 wherein hole transport is provided by a triarylamine repeat unit located within the polymer backbone and electron transport is provided by a conjugated polyfluorene chain within the polymer backbone. Alternatively, the properties of hole transport and electron transport may be provided by repeat units pendant from a conjugated or non-conjugated polymer backbone.

Specific examples of "small molecule" hosts include 4,4'-bis(carbazol-9-yl)biphenyl, known as CBP, and (4,4',4''-tris(carbazol-9-yl)triphenylamine), known as TCTA, disclosed in Ikai et al. (*Appl. Phys. Lett.*, 79 no. 2, 2001, 156); and triarylamine such as tris-4-(N-3-methylphenyl-N-phenyl)phenylamine, known as MTDATA.

Homopolymers and copolymers may be used as hosts, including optionally substituted polyarylenes such as polyfluorenes, polyspirofluorenes, polyindenofluorenes or polyphenylenes as described above with respect to the hole transporting layer.

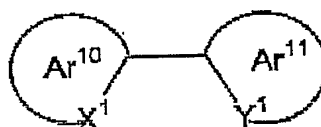
Specific examples of host polymers disclosed in the prior art include poly(vinyl carbazole) disclosed in, for example, Appl. Phys. Lett. 2000, 77(15), 2280; polyfluorenes in Synth. Met. 2001, 116, 379, Phys. Rev. B 2001, 63, 235206 and Appl. Phys. Lett. 2003, 82(7), 1006; poly[4-(N-4-vinylbenzyloxyethyl, N-methylamino)-N-(2,5-di-tert-butylphenyl)naphthalimide] in Adv. Mater. 1999, 11(4), 285; poly(para-phenylenes) in J. Mater. Chem. 2003, 13, 50-55; poly[9,9'-di-n-hexyl-2,7-fluorene-alt-1,4-(2,5-di-n-hexyloxy)phenylene] as a host for both fac-tris(2-phenylpyridine) iridium(III) and 2,3,7,8,12,13, 17,18-octaethyl-21H,23H-porphyrin platinum(II) in J. Chem. Phys. (2003), 118(6), 2853-2864; a random copolymer host of dioctylfluorene and dicyano-benzene in Mat. Res. Symp. Spring Meeting 2003 Book of Abstracts, Heeger, p. 214; and an AB copolymer of a fluorene repeat unit and phenylene repeat unit is disclosed in Mat. Res. Soc. Symp. Proc. 708, 2002, 131.

Suitable phosphorescent dopants are metal complexes based on heavy elements M that induce strong spin-orbit coupling to allow rapid intersystem crossing and emission from triplet states. Suitable heavy metals M include:

- lanthanide metals such as cerium, samarium, europium, terbium, dysprosium, thulium, erbium and neodymium; and
- d-block metals, in particular those in rows 2 and 3 i.e. elements 39 to 48 and 72 to 80, in particular ruthenium, rhodium, palladium, rhenium, osmium, iridium, platinum and gold.

Suitable coordinating groups for the f-block metals include oxygen or nitrogen donor systems such as carboxylic acids, 1,3-diketonates, hydroxy carboxylic acids, Schiff bases including acyl phenols and iminoacyl groups. As is known, luminescent lanthanide metal complexes require sensitizing group(s) which have the triplet excited energy level higher than the first excited state of the metal ion. Emission is from an f-f transition of the metal and so the emission colour is determined by the choice of the metal. The sharp emission is generally narrow, resulting in a pure colour emission useful for display applications.

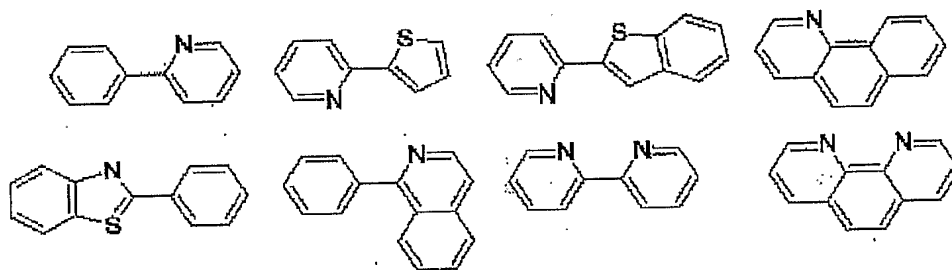
The d-block metals form organometallic complexes with carbon or nitrogen donors such as porphyrin or bidentate ligands of formula (VI):



(VI)

wherein Ar^{10} and Ar^{11} may be the same or different and are independently selected from optionally substituted aryl or heteroaryl; X^1 and Y^1 may be the same or different and are independently selected from carbon or nitrogen; and Ar^{10} and Ar^{11} may be fused together. Ligands wherein X^1 is carbon and Y^1 is nitrogen, or wherein X^1 and Y^1 are both nitrogen are particularly preferred.

Examples of bidentate ligands are illustrated below:



One or both of Ar^{10} and Ar^{11} may carry one or more substituents. Particularly preferred substituents include fluorine or trifluoromethyl which may be used to blue-shift the emission of the complex as disclosed in WO 02/45466, WO 02/44189, US 2002-117662 and US 2002-182441; alkyl or alkoxy groups as disclosed in JP 2002-324679; carbazole which may be used to assist hole transport to the complex when used as an emissive material as disclosed in WO 02/81448; bromine, chlorine or iodine which can serve to functionalise the ligand for attachment of further groups as disclosed in WO 02/68435 and EP 1245659; and dendrons which may be used to obtain or enhance solution processability of the metal complex as disclosed in WO 02/66552.

Other ligands suitable for use with d-block elements include diketonates, in particular acetylacetonate (acac); triarylphosphines and pyridine, each of which may be substituted.

The concentration of the phosphorescent light-emitting dopant in the host material should be such that the film has a high electroluminescent efficiency. If the concentration of the emissive species is too high, quenching of luminescence can occur. A concentration in the range 0.01-49 wt %, more preferably 0.5-10 wt %, most preferably 1-3 wt % is generally appropriate.

The host material and the electroluminescent material may be provided as separate materials as described above. Alternatively, they may be components of the same molecule. For example, a phosphorescent metal complex may be provided as repeat unit, sidechain substituent or end-group of a host polymer as disclosed in, for example, WO 02/31896, WO 03/001616, WO 03/018653 and EP 1245659. Likewise, a "small molecule" host material may be bound directly to a ligand of a phosphorescent metal complex.

The phosphorescent material is typically employed as a dopant within the host material at low (ca. 1-5 %) concentration. It will therefore be appreciated that solubility of the phosphorescent material in a given solvent may be correspondingly low. Alternatively, a phosphorescent material having enhanced solubility may be employed. Such phosphorescent materials include complexes bearing alkyl or alkoxy groups as disclosed in JP 2002-324679; dendrimers as disclosed in WO 02/66552; and phosphorescent complexes provided as repeat units, sidechain substituents or end-groups of soluble polymers as disclosed in WO 02/31896, WO 03/001616, WO 03/018653 and EP 1245659.

Suitable solution processing techniques for deposition of the hole transporting layer and the electroluminescent layer include spin-coating, inkjet printing as disclosed in EP 0880303, laser transfer as described in EP 0851714, flexographic printing, screen printing and doctor blade coating. Provision of such solubilising groups as described above may be particularly advantageous for rendering the phosphorescent material more suitable for solution processing techniques such as inkjet printing.

Following the aforementioned treatment, the hole transporting layer may be entirely insoluble or partially insoluble. In the case where the hole transporting material is substantially free of cross-linkable groups, the degree of insolubility may vary with the duration and / or temperature of heat treatment. Any soluble hole transporting material may be removed from the layer by rinsing in an appropriate solvent in order to leave an entirely insoluble layer, however the invention also encompasses deposition of the electroluminescent layer wherein the first layer has been rendered only partially insoluble and the soluble portion has not been removed. In this instance there will be some blending between the materials of the hole transporting layer and the electroluminescent layer. This may be advantageous in both providing a hole transporting layer and an electroluminescent layer comprising some hole transporting material, thus creating a gradient of increasing concentration of hole transporting material in moving from the cathode towards the anode.

The hole transporting material, the host material and the phosphorescent material according to the invention must be at least partially soluble in a common solvent. Suitable solvents for deposition of the electroluminescent layer and (where solution deposited) the hole transport layer will be apparent to the skilled person. For example, alkyl- or alkoxy- substituted polyarylenes such as polyfluorenes, polyphenylenes and polyindenofluorenes are typically soluble in aromatic hydrocarbons, more preferably mono- or poly- alkylated or alkoxyated benzene such as toluene ethylbenzene and cyclohexylbenzene; polyalkylbenzenes, for example xylene, trimethylbenzene and tetramethylbenzene; alkoxybenzenes such as anisole; and blends thereof. On the other hand, PVK is soluble in solvents such as 1,2-dichloroethane.

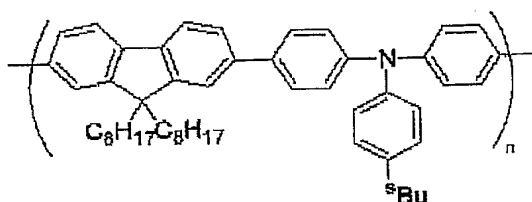
The described treatment has been found to improve efficiency and lifetime of PLEDs. Without wishing to be bound by any theory, the following factors may contribute to these observed increases:

- The first layer may act as a hole transporting, electron blocking and / or exciton blocking layer when located between the anode and the second layer.
- Where a PEDT / PSS layer is used, the insoluble hole transporting layer may prevent ingress of protons from the acidic PSS material into the second layer. This may apply in particular where the polymer comprises basic units such as amines of formulae 1-6 or Het groups described above, such as pyridyl, that are capable of accepting protons.
- The thinness of the insoluble layer, which is achievable in particular when rinsing of the hole transporting layer is employed to remove remaining soluble material, may block electrons from entering the PEDT / PSS hole injection layer and / or ITO anode layer without any detriment to device performance that may result from a thicker layer, such as higher drive voltage.

Examples

General procedure

The invention is exemplified here using the polymer "F8-TFB", illustrated below and disclosed in WO 99/54385, as the hole transporting layer.

**F8-TFB**

The general procedure follows the steps outlined below:

- 1) Depositing PEDT / PSS (available from H C Starck of Leverkusen, Germany as Baytron P ®) onto indium tin oxide supported on a glass substrate (available from Applied Films, Colorado, USA) by spin coating.
- 2) Depositing a hole transporting layer by spin coating F8-TFB from a xylene solution having a concentration of 2 % w / v.
- 3) Heating the hole transporting layer in an inert (nitrogen) environment.
- 4) Optionally spin-rinsing the substrate in xylene to remove any remaining soluble F8-TFB.
- 5) Depositing the electroluminescent layer by spin-coating from xylene solution.
- 6) Depositing over the electroluminescent layer a cathode comprising a first layer of lithium fluoride (4 nm), a second layer calcium (10 nm) and a capping layer of aluminium (>200 nm) as described in WO 00/48258.
- 7) Encapsulation of the device using an airtight metal enclosure available from Saes Getters SpA.

Parameters within this general process may be varied. In particular: the concentration of the hole transporting material may be up to around 3 % w / v. Alternatively, it may be as low as 0.5 % w / v to provide a particularly thin film; the optional heating step may last for any length of time up to around 2 hours; and the optional heating step may be at any temperature up to around 220°C, but preferably above the glass transition temperature of the deposited polymer. As will be apparent to the skilled person, first and / or second polymers, and other device components such as PEDT / PSS, will be subject to thermal degradation if the heating temperature is excessive, and the heating temperature should be selected accordingly.

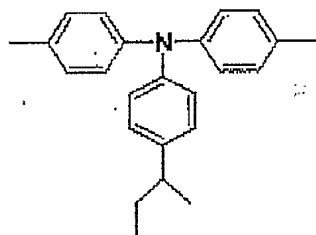
Example 1

The general procedure above was followed using an electroluminescent layer comprising:

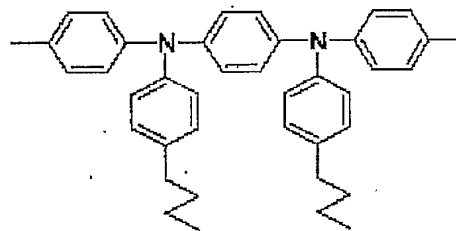
- a host polymer having the composition 70 % 9,9-dioctylfluorene-2,7-diyl, 10 % 9,9-diphenylfluorene-2,7-diyl, 10 % "TFB" repeat unit and 10 % "PFB" repeat unit as disclosed in WO 02/92723 (TFB and PFB repeat units are illustrated below). The polymer was prepared by Suzuki polymerisation as disclosed in, for example, WO 00/53656.
- the red phosphorescent dopant platinum octaethylporphine (PtOEP) as disclosed in *Nature (London)*, 1998, 395, 151.

The host : dopant ratio was 98.8 : 1.2.

The hole transporting layer was subjected to heat treatment at 130°C for 10 minutes prior to deposition of the electroluminescent layer.



"TFB"



"PFB"

Comparative Example 1

For the purpose of comparison, a device was made in accordance with Example 1 except that the hole transporting layer (i.e. steps 2-4 of the above general method) was omitted.

Example 2

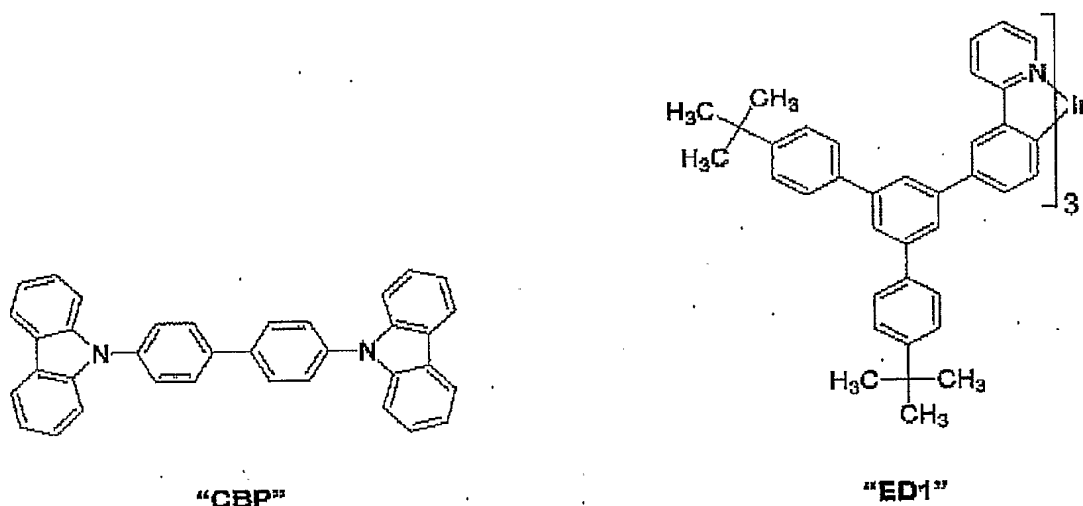
A device was made in accordance with Example 1 except that the host : dopant ratio was 98.0 : 2.0.

Comparative Example 2

For the purpose of comparison, a device was made in accordance with Example 2 except that the hole transporting layer (i.e. steps 2-4 of the above general method) was omitted.

Example 3

A device was made in accordance with Example 1 except that the electroluminescent layer consisted of host material CBP and the green electroluminescent dendrimer dopant ED1, disclosed in WO 02/066552 and illustrated below, deposited by spin-coating from a chloroform solution having a CBP:ED1 ratio of 8:2.



Comparative Example 3

For the purpose of comparison, a device was made in accordance with Example 3 except that the hole transporting layer (i.e. steps 2-4 of the above general method) was omitted.

Table 1 below shows a comparison of quantum efficiency and lifetime of devices according to the invention and the comparative examples. As can be seen, there is a very significant improvement in efficiency upon inclusion of the hole transporting layer for Examples 1 and 2. Furthermore, lifetime is improved for both the red PtOEP emitter (Example 1) and the green ED1 emitter (Example 3).

In contrast to Examples 1 and 2, efficiency falls for Example 3 (ED1 dopant) relative to its Comparative Example. Without wishing to be bound by any theory, this is believed to be due to quenching of luminance by hole transporting layer 4 in Example 3 as a result of the bandgap of this material being insufficiently wide as compared to the bandgap of the electroluminescent material. It is believed that the increase in lifetime of Example 3 may be achieved whilst

retaining or exceeding the efficiency of the Comparative Example by use of an appropriate wider bandgap material (i.e. an exciton blocking layer as hereinbefore described).

TABLE 1

Device	Lifetime*	Quantum efficiency at 100 cd / m ²	Maximum quantum efficiency
Comparative Example 1	3852 ¹	1.88	2.26
Example 1	9592 ²	3.25	3.52
Comparative Example 2		1.92	2.22
Example 2		2.98	3.11
Comparative Example 3	670	6.4	6.4
Example 3	1014	3.0	4.2

*"Lifetime" in Table 1 is the time taken for luminance of the device to halve when driven at a constant current from a starting luminance of 400 cd/m².

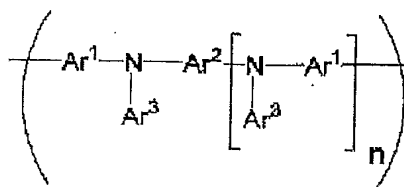
¹ Average of 2 devices measured

² Average of 3 devices measured

Although the present invention has been described in terms of specific exemplary embodiments, it will be appreciated that various modifications, alterations and / or combinations of features disclosed herein will be apparent to those skilled in the art without departing from the spirit and scope of the invention as set forth in the following claims.

Claims

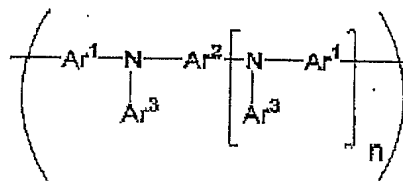
1. A method of forming an organic light emitting diode comprising the steps of:
 - providing a substrate comprising a first electrode for injection of charge carriers of a first type
 - forming a charge transporting layer by depositing onto the substrate a charge transporting material for transporting charge carriers of the first type, the charge transporting material being soluble in a solvent;
 - treatment of the charge transporting layer to render it insoluble in the solvent;
 - forming an electroluminescent layer by depositing onto the charge transporting layer a composition comprising the solvent, a phosphorescent material and a host material; and
 - depositing onto the electroluminescent layer a second electrode for injection of charge carriers of a second type.
2. A method according to claim 1 wherein the first electrode is an anode; the second electrode is a cathode; the charge carriers of the first type are holes; and the charge carriers of the second type are electrons.
3. A method according to claim 1 or 2 wherein the charge transporting material comprises a cross-linkable material and the treatment comprises subjecting the charge transporting layer to heat or electromagnetic radiation in order to cross-link the charge transporting material
4. A method according to claim 1 or 2 wherein the charge transporting layer is substantially free of cross-linkable groups and the treatment comprises subjecting the charge transporting layer to heat.
5. A method according to any one of claims 1-4 wherein the charge transporting material is a polymer.
6. A method according to claim 5 wherein the polymer comprises an optionally substituted triarylamine repeat unit.
7. A method according to claim 6 wherein the triarylamine repeat unit comprises an optionally substituted repeat unit of formula (I):



(I)

wherein each Ar¹, Ar² and Ar³ is the same or different and independently represents optionally substituted aryl; and n is 0 or 1.

8. A method according to any one of claims 5-7 wherein the polymer comprises a repeat unit selected from optionally substituted fluorene, indenofluorene, spirofluorene and phenylene.
9. A method according to any preceding claim wherein the phosphorescent material is a metal complex.
10. A method according to any preceding claim wherein the host material is a host polymer.
11. A method according to claim 10 wherein the host polymer comprises a repeat unit as defined in claim 7 or claim 8.
12. An organic light emitting diode obtainable by the method according to any preceding claim.
13. An organic light emitting diode comprising, in sequence, an anode; a hole transporting layer; an electroluminescent layer comprising a phosphorescent material and a host material; and a cathode, wherein the hole transporting layer is a polymer comprising an optionally substituted repeat unit of formula (I):



(I)

wherein each Ar¹, Ar² and Ar³ is the same or different and independently represents optionally substituted aryl; and n is 0 or 1.

14. An organic light emitting diode according to claim 13 wherein the polymer comprises a repeat unit selected from optionally substituted fluorene, indenofluorene, spirofluorene and phenylene.
15. An organic light emitting diode according to claim 13 or 14 wherein a hole injecting layer comprising a conductive organic material is located between the anode and the hole transporting layer.
16. An organic light emitting diode according to any one of claims 13-15 wherein the phosphorescent material is a metal complex.